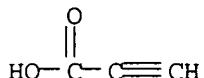


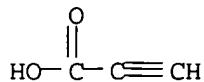
RN 471-25-0 REGISTRY  
ED Entered STN: 16 Nov 1984  
CN 2-Propynoic acid (9CI) (CA INDEX NAME)  
OTHER CA INDEX NAMES:  
CN Propiolic acid (6CI, 8CI)  
OTHER NAMES:  
CN 2-Propyne-1-carboxylic acid  
CN Acetylenecarboxylic acid  
CN Acetylenemonocarboxylic acid  
CN Carboxyacetylene  
CN NSC 16152  
CN Propargylic acid  
CN Propynoic acid  
FS 3D CONCORD  
MF C3 H2 O2  
CI COM  
LC STN Files: AGRICOLA, AQUIRE, BEILSTEIN\*, BIOBUSINESS, BIOSIS, CA, CAOLD, CAPLUS, CASREACT, CHEMCATS, CHEMINFORMRX, CHEMLIST, CHEMSAFE, CSCHEM, DDFU, DETHERM\*, DRUGU, EMBASE, GMELIN\*, HODOC\*, IFICDB, IFIPAT, IFIUDB, MEDLINE, MRCK\*, RTECS\*, SPECINFO, SYNTHLINE, TOXCENTER, USPAT2, USPATFULL  
(\*File contains numerically searchable property data)  
Other Sources: EINECS\*\*, NDSL\*\*, TSCA\*\*  
(\*\*Enter CHEMLIST File for up-to-date regulatory information)



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

935 REFERENCES IN FILE CA (1907 TO DATE)  
62 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
935 REFERENCES IN FILE CAPLUS (1907 TO DATE)

IT 471-25-0P  
RL: IMF (Industrial manufacture); PREP (Preparation)  
(production of, electrochem., extraction in relation to)  
RN 471-25-0 CAPLUS  
CN 2-Propynoic acid (9CI) (CA INDEX NAME)



=> d his

(FILE 'HOME' ENTERED AT 11:16:12 ON 21 APR 2005)

FILE 'REGISTRY' ENTERED AT 11:16:21 ON 21 APR 2005

L1 1 S PROPIOLIC ACID/CN  
L2 0 S ACETYLENE DICARBOXYLIC ACID/CN  
L3 108 S (ACETYLENE(3A)DICARBOXYLIC)  
L4 107 S L3 AND (DICARBOXYLIC(W)ACID)

FILE 'CAPLUS' ENTERED AT 11:24:36 ON 21 APR 2005

L5 36 S 142-45-0P  
L6 1 S L5 AND (H2O2 OR ?PEROXID?)  
L7 2 S L5 AND (HYPOCHLOR? OR HYPOHAL? OR HYPOIOD? OR HYPOBROM?)  
L8 2 S L5 AND NITROXY?  
L9 60 S 471-25-0P  
L10 1 S L9 AND (?PEROXID? OR H2O2)

=> s 19 and (hypohal? or hypochlo? or hypobrom? or hypoiodo?)

1351 HYPOHAL?  
27642 HYPOCHLO?  
2899 HYPOBROM?  
322 HYPOIODO?

L11 3 L9 AND (HYPOHAL? OR HYPOCHLO? OR HYPOBROM? OR HYPOIODO?)

=> d bib abs hit 1-3

L11 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN  
AN 2004:159015 CAPLUS  
DN 140:199022  
TI Procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcohols with **hypohalites** in the presence of a nitroxyl compound

IN Stohrer, Juergen; Fritz-Langhals, Elke; Bruenninghaus, Christian  
PA Consortium fuer Elektrochemische Industrie G.m.b.H., Germany  
SO Ger., 11 pp.  
CODEN: GWXXAW

DT Patent  
LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 10244633	B3	20040226	DE 2002-10244633	20020925
	EP 1403240	A1	20040331	EP 2003-20442	20030911
	EP 1403240	B1	20040721		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	AT 271533	E	20040815	AT 2003-20442	20030911

ES 2222450	T3	20050201	ES 2003-3020442	20030911
US 2004059154	A1	20040325	US 2003-667810	20030922
JP 2004115519	A2	20040415	JP 2003-331417	20030924
PRAI DE 2002-10244633	A	20020925		
OS CASREACT 140:199022				

AB Alkyne carboxylic acids (e.g., propargylic acid) are prepared in high yield and selectivity by the oxidation of an alkynyl alc. (e.g., propargylic alc.) with a **hypohalite** (e.g., sodium **hypochlorite**) in the presence of a nitroxyl compound (e.g., 4-hydroxy-TEMPO) at a pH value >7 by continuous addition of the alkynyl alc. and the **hypohalogenite** to the reaction mixture

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

TI Procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcohols with **hypohalites** in the presence of a nitroxyl compound

AB Alkyne carboxylic acids (e.g., propargylic acid) are prepared in high yield and selectivity by the oxidation of an alkynyl alc. (e.g., propargylic alc.) with a **hypohalite** (e.g., sodium **hypochlorite**) in the presence of a nitroxyl compound (e.g., 4-hydroxy-TEMPO) at a pH value >7 by continuous addition of the alkynyl alc. and the **hypohalogenite** to the reaction mixture

ST alkyne carboxylic acid manuf alkynyl alc oxidn **hypohalite**  
nitroxyl compd; propyne carboxylic acid manuf propargylic alc oxidn  
**hypohalite** nitroxyl compd

IT Buffers  
(in a procedure for the production of alkyne carboxylic acids by the oxidation

of alkynyl alcs. with **hypohalites** in the presence of a  
nitroxyl compound)

IT Oxidation catalysts  
(liquid-phase, phase-transfer; in a procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with  
**hypohalites** in the presence of a nitroxyl compound)

IT Oxidation  
(liquid-phase; procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a  
nitroxyl compound)

IT Hypochlorites  
**Hypohalites**

RL: RCT (Reactant); RACT (Reactant or reagent)  
(oxidants; procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a  
nitroxyl compound)

IT Nitroxides  
RL: CAT (Catalyst use); USES (Uses)  
(procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl  
compound)

IT Alcohols, reactions  
RL: RCT (Reactant); RACT (Reactant or reagent)  
(propargyl; procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a  
nitroxyl compound)

IT Carboxylic acids, preparation  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(unsatd., alkyne carboxylic acids; procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with  
**hypohalites** in the presence of a nitroxyl compound)

IT 471-34-1, Calcium carbonate, reactions  
RL: RGT (Reagent); RACT (Reactant or reagent)  
(base; in a procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a

nitroxyl compound)

IT 2226-96-2, 4-Hydroxy-TEMPO  
 RL: CAT (Catalyst use); USES (Uses)  
 (in a procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT 7681-52-9, Sodium **hypochlorite**  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (oxidant; procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT 14691-89-5, 4-Acetamido-TEMPO  
 RL: CAT (Catalyst use); USES (Uses)  
 (procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT 107-19-7, Propargyl alcohol 110-65-6, 2-Butyne-1,4-diol 764-01-2, 2-Butyn-1-ol  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT 142-45-0P, Acetylenedicarboxylic acid 471-25-0P, Propargylic acid 590-93-2P, 2-Butynoic acid  
 RL: SPN (Synthetic preparation); PREP (Preparation)  
 (procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

L11 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2003:652130 CAPLUS

DN 139:181969

TI Process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols

IN Stohrer, Juergen; Fritz-Langhals, Elke; Brueninghaus, Christian; Stauch, Dagmar

PA Consortium Fuer Elektrochemische Industrie G.m.b.H., Germany

SO Eur. Pat. Appl., 10 pp.

CODEN: EPXXDW

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1336599	A1	20030820	EP 2003-2103	20030130
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	DE 10206619	A1	20031009	DE 2002-10206619	20020215
	DE 10206619	B4	20040325		
	US 2003158439	A1	20030821	US 2003-365887	20030213
PRAI	DE 2002-10206619	A	20020215		

OS CASREACT 139:181969

AB Alkynoic acids (e.g., propynoic acid) and alkynoic acid esters of alkynols (e.g., 2-propyn-1-yl propynoate) are prepared in high yield and selectivity via the oxidation of alkynols (e.g., propargyl alc.) in the presence of 1-10 mol-equivalent of a **hypohalogenite** (e.g., sodium **hypochlorite**) and in the presence of a nitroxy compds. (e.g., TEMPO) at a pH of <7.

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

AB Alkynoic acids (e.g., propynoic acid) and alkynoic acid esters of alkynols (e.g., 2-propyn-1-yl propynoate) are prepared in high yield and selectivity

via the oxidation of alkynols (e.g., propargyl alc.) in the presence of 1-10 mol-equivalent of a **hypohalogenite** (e.g., sodium **hypochlorite**) and in the presence of a nitroxy compds. (e.g., TEMPO) at a pH of <7.

IT Oxidizing agents

(**hypohalogenites**; process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols using)

IT **Hypohalites**

RL: RGT (Reagent); RACT (Reactant or reagent)

(process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols using)

IT 7681-52-9, Sodium **hypochlorite**

RL: RGT (Reagent); RACT (Reactant or reagent)

(oxidant; process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols)

IT 471-25-0P, Propynoic acid

RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols)

L11 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1973:526292 CAPLUS

DN 79:126292

TI Synthesis and reactions of acetylenic heterocyclic acids

AU Azerbaev, I. N.; Kurmangazieva, Zh. M.; Nurgalieva, A. N.; Khairova, F. Kh.; Shunkarov, U. Sh.; Yagudeev, T. A.; Leonov, I. D.; Sarbaev, T. G.

CS USSR

SO Khim. Atsetilena Tekhnol. Karbida Kal'tsiya (1972) 131-6

From: Ref. Zh., Khim. 1973, Abstr. No. 8Zh321

DT Journal

LA Russian

GI For diagram(s), see printed CA Issue.

AB Reaction of aqueous KOBr on I or II ( $\beta$ - and  $\gamma$ -isomers) (R = C.tplbond.CH) in diglyme, dioxane, THF, ethylene glycol, or diethylene glycol at -10° gave 80% I or II (R = C.tplbond.CBr) (Ia, IIa), which were hydrogenated over Raney Ni or hydrated to the resp. I and II (R = CH<sub>2</sub>CH<sub>2</sub>Br.COCH<sub>2</sub>Br). Iotsich reaction (Mg, CO<sub>2</sub>) of Ia or IIa 8-10 hr at -20° gave I and II (R = C.tplbond.CCO<sub>2</sub>H) (Ib, IIb), which with MeOH-Wolfatite II give the Me esters, and which were hydrogenated over Raney Ni to give I and II (R = CH<sub>2</sub>CH<sub>2</sub>CO<sub>2</sub>H). Hydration of Ib and IIb gave, resp., III and IV. Oxidation of HOCMe<sub>2</sub>C.tplbond.CCH:CH<sub>2</sub> with ACOOH gave HOCMe<sub>2</sub>C.tplbond.CCO<sub>2</sub>H, which by reverse Favorskii reaction in the presence of KOH gave HC.tplbond.CCO<sub>2</sub>H, condensation of which with Ia or IIa in the presence of CuCl gave I or II (R = C.tplbond.CC.tplbond.CCO<sub>2</sub>H).

IT 74-86-2DP, Ethyne, heterocyclic 471-25-0P 39595-78-3P

42414-47-1P 50288-79-4P 50288-81-8P 50288-83-0P 50624-16-3P

50624-17-4P 50624-18-5P 50624-19-6P 50624-20-9P 50624-21-0P

50624-22-1P 50624-23-2P 50624-24-3P 50624-25-4P 50624-26-5P

50624-27-6P

RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation of)

IT 15775-89-0 19973-20-7

RL: RCT (Reactant); RACT (Reactant or reagent)  
(reaction of, with potassium **hypobromite**)

AN 1961:130815 CAPLUS

DN 55:130815

OREF 55:24569f-g

TI Halogenated acetylenic alcohols

IN Russell, James P.; Vitcha, James F.

PA Air Reduction Co., Inc.

DT Patent

LA Unavailable

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2989568		19610620	US	

L23 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2003:150421 CAPLUS  
 DN 138:172129  
 TI Making carboxylated cellulose fibers and paper products  
 IN Jewell, Richard A.; Komen, Joseph Lincoln; Su, Bing; Weerawarna, S.  
 Ananda; Li, Yong  
 PA Weyerhaeuser Company, USA  
 SO U.S., 23 pp., Cont.-in-part of U.S. 6,379,494.  
 CODEN: USXXAM  
 DT Patent  
 LA English  
 FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 6524348	B1	20030225	US 2000-641276	20000817
	US 6379494	B1	20020430	US 1999-418909	19991015
PRAI	US 1999-272137	B2	19990319		
	US 1999-418909	A2	19991015		

OS MARPAT 138:172129

RE.CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

AB The title method of making carboxylated cellulose fibers whose fiber strength and d.p. is not significantly sacrificed comprises oxidation and **stabilized** stages. The title method involves the use of cyclic **nitroxide** free radical compds. as a primary oxidant and a **hypohalite** salt as a secondary oxidant in an aqueous environment. Preferably the oxidized cellulose is then **stabilized** against D.P. loss in alkaline environments and color reversion with a reducing agent such as Na borohydride. Alternatively it may be treated with an tertiary oxidant such as Na chlorite. The method results in a high percentage of carboxyl groups located at the fiber surface. The product is especially useful as a papermaking fiber where it contributes strength and has a higher attraction for cationic additives. The product is also useful as an additive to recycled fiber to increase strength. The method can be used to improve properties of either virgin or recycled fiber. It does not require high  $\alpha$ -cellulose fiber but is suitable for regular market pulps.

L23 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2001:300943 CAPLUS  
 DN 134:312682  
 TI Method of making carboxylated cellulose fibers and products  
 IN Jewell, Richard A.; Komen, Joseph Lincoln; Su, Bing; Weerawarna, S.  
 Ananda; Li, Yong  
 PA Weyerhaeuser Company, USA  
 SO PCT Int. Appl., 52 pp.  
 CODEN: PIXXD2

DT Patent  
 LA English  
 FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001029309	A1	20010426	WO 2000-US27837	20001006
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,				

CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG  
US 6379494 B1 20020430 US 1999-418909 19991015  
CA 2384701 AA 20010426 CA 2000-2384701 20001006  
EP 1238142 A1 20020911 EP 2000-970682 20001006  
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,  
IE, SI, LT, LV, FI, RO, MK, CY, AL  
JP 2003512540 T2 20030402 JP 2001-532283 20001006  
PRAI US 1999-418909 A 19991015  
US 1999-272137 A2 19990319  
WO 2000-US27837 W 20001006  
OS MARPAT 134:312682

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

AB A method of making highly carboxylated cellulose fibers whose fiber strength and d.p. is not significantly sacrificed comprises (1) oxidizing the cellulose fiber (kraft pulp) with a cyclic **nitroxide** free radical compound as a primary oxidant and a **hypohalite** salt as a secondary oxidant under aqueous alkaline conditions; and (2) treating the oxidized cellulose against d.p. loss in aqueous suspension with a **stabilizing** agent selected from the group consisting of reducing agent and tertiary oxidizing agent. The product is especially useful as a papermaking fiber where it contributes strength and has a higher attraction for cationic additives, and it is also useful as an additive to recycled fiber to increase strength.

RN 142-45-0 REGISTRY

ED Entered STN: 16 Nov 1984

CN 2-Butynedioic acid (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Acetylenedicarboxylic acid (6CI, 8CI)

OTHER NAMES:

CN 2-Butyne-1,4-dioic acid

CN Butynedioic acid

CN NSC 1903

CN NSC 631597

FS 3D CONCORD

MF C4 H2 O4

CI COM

LC STN Files: BEILSTEIN\*, BIOSIS, CA, CAOLD, CAPLUS, CASREACT, CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, DETHERM\*, EMBASE, GMELIN\*, HODOC\*, IFICDB, IFIPAT, IFIUDB, MEDLINE, MSDS-OHS, SPECINFO, TOXCENTER, USPAT2, USPATFULL

(\*File contains numerically searchable property data)

Other Sources: EINECS\*\*, NDSL\*\*, TSCA\*\*

(\*\*Enter CHEMLIST File for up-to-date regulatory information)



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

562 REFERENCES IN FILE CA (1907 TO DATE)

71 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

564 REFERENCES IN FILE CAPLUS (1907 TO DATE)

48 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

AN 1973:525868 CAPLUS  
DN 79:125868  
TI Acetylenedicarboxylic acid  
IN Vereshchagin, L. I.; Gavrilov, L. D.  
PA Irkutsk State University  
SO U.S.S.R.  
From: Otkrytiya, Izobret., Prom. Obraztsy, Tovarnye Znaki 1973, 50(29),  
88.

CODEN: URXXAF

DT Patent

LA Russian

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	SU 389075	T	19730705	SU 1971-1710938	19711101
PRAI	SU 1971-1710938	A	19711101		
AB	HO2CC.tplbond.CCO2H was prepared by oxidation of HOCH2CH.tplbond.CHCH2OH with nickel <b>peroxide</b> in aqueous base.				
ST	acetylenedicarboxylic acid; butenediol nickel <b>peroxide</b> oxidn				
IT	Oxidation (of butenediol by nickel <b>peroxide</b> , acetylenedicarboxylic acid from)				
IT	<b>142-45-0P</b> RL: PREP (Preparation) (by oxidation of butenediol)				
IT	110-64-5 RL: RCT (Reactant); RACT (Reactant or reagent) (oxidation of, with nickel <b>peroxide</b> )				

RN 142-45-0 REGISTRY

ED Entered STN: 16 Nov 1984

CN 2-Butynedioic acid (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Acetylenedicarboxylic acid (6CI, 8CI)

OTHER NAMES:

CN 2-Butyne-1,4-dioic acid

CN Butynedioic acid

CN NSC 1903

CN NSC 631597

FS 3D CONCORD

MF C4 H2 O4

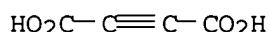
CI COM

LC STN Files: BEILSTEIN\*, BIOSIS, CA, CAOLD, CAPLUS, CASREACT, CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, DETHERM\*, EMBASE, GMELIN\*, HODOC\*, IFICDB, IFIPAT, IFIUDB, MEDLINE, MSDS-OHS, SPECINFO, TOXCENTER, USPAT2, USPATFULL

(\*File contains numerically searchable property data)

Other Sources: EINECS\*\*, NDSL\*\*, TSCA\*\*

(\*\*Enter CHEMLIST File for up-to-date regulatory information)



=> s 142-45-0p  
L5 36 142-45-0P

=> s 15 and (h2o2 or ?peroxid?)  
143460 H2O2  
377216 ?PEROXID?  
L6 1 L5 AND (H2O2 OR ?PEROXID?)

=> d bib hit

L6 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN  
AN 1973:525868 CAPLUS  
DN 79:125868  
TI Acetylenedicarboxylic acid  
IN Vereshchagin, L. I.; Gavrilov, L. D.  
PA Irkutsk State University  
SO U.S.S.R.  
From: Otkrytiya, Izobret., Prom. Obraztsy, Tovarnye Znaki 1973, 50(29), 88.

CODEN: URXXAF

DT Patent

LA Russian

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	SU 389075	T	19730705	SU 1971-1710938	19711101
PRAI	SU 1971-1710938	A	19711101		

AB HO<sub>2</sub>CC.tplbond.CCO<sub>2</sub>H was prepared by oxidation of HOCH<sub>2</sub>CH.tplbond.CHCH<sub>2</sub>OH with nickel **peroxide** in aqueous base.

ST acetylenedicarboxylic acid; butenediol nickel **peroxide** oxidn

IT Oxidation

(of butenediol by nickel **peroxide**, acetylenedicarboxylic acid from)

IT 142-45-0P

RL: PREP (Preparation)  
(by oxidation of butenediol)

IT 110-64-5

RL: RCT (Reactant); RACT (Reactant or reagent)  
(oxidation of, with nickel **peroxide**)

=> s 15 and (hypochlor? or hypohal? or hypoiod? or hypobrom?)

27623 HYPOCHLOR?

1351 HYPOHAL?

1147 HYPOIOD?

2899 HYPOBROM?

L7 2 L5 AND (HYPOCHLOR? OR HYPOHAL? OR HYPOIOD? OR HYPOBROM?)

=> d bib abs hit 1-2

L7 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:159015 CAPLUS

DN 140:199022

TI Procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcohols with **hypohalites** in the presence of a nitroxyl compound

IN Stohrer, Juergen; Fritz-Langhals, Elke; Bruenninghaus, Christian

PA Consortium fuer Elektrochemische Industrie G.m.b.H., Germany

SO Ger., 11 pp.

CODEN: GWXXAW

DT Patent

LA German

FAN. CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 10244633	B3	20040226	DE 2002-10244633	20020925
	EP 1403240	A1	20040331	EP 2003-20442	20030911
	EP 1403240	B1	20040721		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	AT 271533	E	20040815	AT 2003-20442	20030911
	ES 2222450	T3	20050201	ES 2003-3020442	20030911
	US 2004059154	A1	20040325	US 2003-667810	20030922
	JP 2004115519	A2	20040415	JP 2003-331417	20030924
PRAI	DE 2002-10244633	A	20020925		
OS	CASREACT 140:199022				

AB Alkyne carboxylic acids (e.g., propargylic acid) are prepared in high yield and selectivity by the oxidation of an alkynyl alc. (e.g., propargylic alc.) with a **hypohalite** (e.g., sodium **hypochlorite**) in the presence of a nitroxyl compound (e.g., 4-hydroxy-TEMPO) at a pH value >7 by continuous addition of the alkynyl alc. and the **hypohalogenite** to the reaction mixture

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

TI Procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcohols with **hypohalites** in the presence of a nitroxyl compound

AB Alkyne carboxylic acids (e.g., propargylic acid) are prepared in high yield and selectivity by the oxidation of an alkynyl alc. (e.g., propargylic alc.) with a **hypohalite** (e.g., sodium **hypochlorite**) in the presence of a nitroxyl compound (e.g., 4-hydroxy-TEMPO) at a pH value >7 by continuous addition of the alkynyl alc. and the **hypohalogenite** to the reaction mixture

ST alkyne carboxylic acid manuf alkynyl alc oxidn **hypohalite** nitroxyl compd; propyne carboxylic acid manuf propargylic alc oxidn **hypohalite** nitroxyl compd

IT Buffers  
(in a procedure for the production of alkyne carboxylic acids by the oxidation

of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT Oxidation catalysts  
(liquid-phase, phase-transfer; in a procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT Oxidation  
(liquid-phase; procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT **Hypochlorites**  
**Hypohalites**

RL: RCT (Reactant); RACT (Reactant or reagent)  
(oxidants; procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT Nitroxides

RL: CAT (Catalyst use); USES (Uses)  
(procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

IT Alcohols, reactions

RL: RCT (Reactant); RACT (Reactant or reagent)  
(propargyl; procedure for the production of alkyne carboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a

nitroxyl compound)  
 IT Carboxylic acids, preparation  
 RL: SPN (Synthetic preparation); PREP (Preparation)  
 (unsatd., alkynecarboxylic acids; procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)  
 IT 471-34-1, Calcium carbonate, reactions  
 RL: RGT (Reagent); RACT (Reactant or reagent)  
 (base; in a procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)  
 IT 2226-96-2, 4-Hydroxy-TEMPO  
 RL: CAT (Catalyst use); USES (Uses)  
 (in a procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)  
 IT 7681-52-9, Sodium **hypochlorite**  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (oxidant; procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)  
 IT 14691-89-5, 4-Acetamido-TEMPO  
 RL: CAT (Catalyst use); USES (Uses)  
 (procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)  
 IT 107-19-7, Propargyl alcohol 110-65-6, 2-Butyne-1,4-diol 764-01-2,  
 2-Butyn-1-ol  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)  
 IT 142-45-0P, Acetylenedicarboxylic acid 471-25-0P, Propargylic acid 590-93-2P, 2-Butynoic acid  
 RL: SPN (Synthetic preparation); PREP (Preparation)  
 (procedure for the production of alkynecarboxylic acids by the oxidation of alkynyl alcs. with **hypohalites** in the presence of a nitroxyl compound)

L7 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2003:652130 CAPLUS  
 DN 139:181969  
 TI Process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols  
 IN Stohrer, Juergen; Fritz-Langhals, Elke; Brueninghaus, Christian; Stauch, Dagmar  
 PA Consortium Fuer Elektrochemische Industrie G.m.b.H., Germany  
 SO Eur. Pat. Appl., 10 pp.  
 CODEN: EPXXDW  
 DT Patent  
 LA German  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1336599	A1	20030820	EP 2003-2103	20030130
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	DE 10206619	A1	20031009	DE 2002-10206619	20020215
	DE 10206619	B4	20040325		
	US 2003158439	A1	20030821	US 2003-365887	20030213
PRAI	DE 2002-10206619	A	20020215		
OS	CASREACT 139:181969				

AB Alkynoic acids (e.g., propynoic acid) and alkynoic acid esters of alkynols (e.g., 2-propyn-1-yl propynoate) are prepared in high yield and selectivity via the oxidation of alkynols (e.g., propargyl alc.) in the presence of 1-10 mol-equivalent of a **hypohalogenite** (e.g., sodium **hypochlorite**) and in the presence of a nitroxy compds. (e.g., TEMPO) at a pH of <7.

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

AB Alkynoic acids (e.g., propynoic acid) and alkynoic acid esters of alkynols (e.g., 2-propyn-1-yl propynoate) are prepared in high yield and selectivity via the oxidation of alkynols (e.g., propargyl alc.) in the presence of 1-10 mol-equivalent of a **hypohalogenite** (e.g., sodium **hypochlorite**) and in the presence of a nitroxy compds. (e.g., TEMPO) at a pH of <7.

IT Oxidizing agents

(**hypohalogenites**; process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols using)

IT **Hypohalites**

RL: RGT (Reagent); RACT (Reactant or reagent)

(process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols using)

IT 7681-52-9, Sodium **hypochlorite**

RL: RGT (Reagent); RACT (Reactant or reagent)

(oxidant; process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols)

IT **142-45-0P**, 2-Butynedioic acid 2345-51-9P, 3-Butynoic acid

4383-39-5P

RL: SPN (Synthetic preparation); PREP (Preparation)

(process for the preparation of alkynoic acids and alkynoic acid esters of alkynols via the oxidation of alkynols)